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In a multivalley semiconductor, an electric field  $E_x$  gives rise to transverse currents of electrons belonging to different valleys

$$j_y^{(\alpha)} = -\mu_{yx}^{(\alpha)} n^{(\alpha)} E_x \quad (1)$$

( $\mu_{ik}^{(\alpha)}$  - mobility tensor in  $\alpha$ -th valley). If the semiconductor is homogeneous in the  $y$  direction then  $n^{(\alpha)} = n/m$  everywhere except in a thin surface region ( $n$  - total electron density,  $m$  - number of equivalent valleys), so that in a crystal with cubic symmetry

$$\sum_{\alpha=1}^m \mu_{yx}^{(\alpha)} = 0$$

the current in the  $y$  direction equals 0. In the presence of a gradient  $dn/dy$ , the continuity of the current  $j_y^{(\alpha)}$  is violated, leading to the appearance of an intervalley redistribution

$$n^{(\alpha)} - \frac{n}{m} = -\tau \frac{dj_y^{(\alpha)}}{dy} \approx \mu_{yx}^{(\alpha)} \tau \frac{E_x}{m} \frac{dn}{dy}, \quad (2)$$

where  $\tau$  is the intervalley relaxation time. Deviation from the intervalley equilibrium (2) gives rise to anisotropy of the conductivity of the semiconductor, so that the total electron current in the  $y$  direction now consists of the diffusion current, connected with the gradient  $dn/dy$ , the drift current in the field  $E_y$ , and the anisotropic current in the field  $E_x$ . Since the latter is proportional to the  $dn/dy$ , a change takes place in the effective coefficient of diffusion in the  $y$  direction

$$j_y = -D_n^* \frac{dn}{dy} - \mu_n n E_y, \quad (3)$$

where

$$D_n^* = D_n (1 + \gamma^2), \quad (4)$$

$D_n = (kT/e)\mu_n$  is the average (isotropic) electron mobility, and  $\gamma$  is the dimensionless electric field  $E_x$ :

$$\gamma^2 = \frac{1}{m} \sum_{\alpha=1}^m \left( \frac{\mu_{yx}^{(\alpha)}}{D_n} E_x L \right)^2, \quad L = \sqrt{D_n \tau}. \quad (5)$$

Formulas (3) - (5) were obtained assuming a small redistribution, meaning smooth inhomogeneity

$$\gamma L \ll L_p, \quad (6)$$

where  $L_p$  is the characteristic length of the inhomogeneity. We note that in the opposite case - near a sharp inhomogeneity - phenomena similar to those considered in [1, 2] come into play.

We now derive some of the consequences of the growth of the diffusion coefficient.

### 1. Monopolar Semiconductor

In this case the carrier-concentration gradient can be obtained by homogeneous doping. The condition that there be no transverse current ( $j_y = 0$ ) leads to the appearance of a transverse volume emf

$$V_y = - \int_{y_1}^{y_2} E_y dy = - \frac{kT}{e} \ln \frac{n_2}{n_1} = \gamma^2 \frac{kT}{e} \ln \frac{n_2}{n_1}. \quad (7)$$

The emf  $V_y$  is strictly proportional to the square of  $E_x$ ; its sign is determined by the sign of the concentration gradient, and its magnitude is proportional to  $\tau$  (and can apparently be used for its measurement).

In the derivation of formula (7) it was assumed that  $\tau$  does not depend on  $y$ . At low temperatures and high impurity concentrations (see [3, 4]),  $\tau$  depends on the concentration and on the degree of donor ionization. In the simplest case, when all the donors are ionized and quasineutrality takes place, it can be assumed that  $\tau = (a + bn)^{-1}$ ; then

$$V_y = \gamma_0^2 \frac{kT}{e} \ln \left[ \left( \frac{n_2}{n_1} \right) \left( \frac{a + bn_1}{a + bn_2} \right) \right], \quad (8)$$

where  $\gamma_0$  is the value of  $\gamma$  at  $\tau = a^{-1}$ .

### 2. Bipolar Nonequilibrium Semiconductor

The inhomogeneous distribution of the electrons in the semiconductor can be produced by photogeneration that is inhomogeneous in the  $y$  direction (or else by homogeneous generation but as a result of the influence of surface recombination). The coefficient of bipolar diffusion in the  $y$  direction is then equal to

$$D = \frac{D_p (D_n^* p + D_n n)}{D_n n + D_p p}, \quad (9)$$

i.e., it remains equal to  $D_p$  in an n-type semiconductor and becomes equal to  $D_n^*$  in a p-type semiconductor. When  $p = n$  we get

$$D = 2D_p D_n / (D_p + D_n) [1 + (\gamma^2/2)].$$

The growth of the diffusion coefficient in the  $y$  direction with increasing field  $E_x$  leads to a "stretching" of the diffusion lengths of the carriers with respect to recombination

$$\left. \begin{aligned} L_p &= L_p^0 (1 + \gamma^2)^{1/2} \text{ for } p \gg n, \\ L_p &= L_p^0 [1 + (\gamma^2/2)]^{1/2} \text{ for } p = n. \end{aligned} \right\} \quad (10)$$

(We note that the criterion (6) is always satisfied here if  $L_p^0 \gg L$ , which is usually the case by a large margin).

The dependence  $L_p$  on the drawing field  $E_x$  greatly influences the field dependence of the photocurrent (cf., e.g., [5]).

In conclusion we note that inasmuch as in germanium the length of the intervalley scattering  $L$  remains essentially larger than the cooling length in almost all situations (the only exception being cases of very high densities and extremely low temperatures, the condition  $\gamma > 1$ , which is necessary for an appreciable stretching of  $L_p$ , is still satisfied in non-heating fields. However, at low temperatures, the intervalley scattering time  $\tau$ , determined by the scattering with emission of intervalley phonons, may turn out to be sensitive to small degrees of heating, so that in general  $\tau = \tau(E_x)$ . Finally, in the bipolar case, the intervalley scattering may proceed via an intermediate exciton state (similar to the proposed intervalley scattering via a bound state by donors in [3, 4]). Then  $\tau$  will depend on  $p$  and will depend on  $y$ , together with the concentration, at high illumination levels ( $p \approx n$ ).

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